directly to the OS OMB Desk Officer all comments must be faxed to OMB at 202–395–6974.

Proposed Project: Institutional Review Board/Independent Ethics Committee Forms Modification–OMB No. 0990–0279—Office for Human Research Protections.

Abstract: The Office for Human Research Protections (OHRP) is requesting a modification to the current Institutional Review Board (IRB) Independent Ethics Committee (IEC) Registration Form designed to provide a simplified procedure for institutions engaged in Department of Health and Human Services (HHS) conducted or supported research to satisfy the assurance requirements of Section 491(a) of the Public Health Service Act and HHS regulations for the protection of human subjects at 45 CFR 46.103. The form is being modified to be consistent with IRB-Registration requirements that are included in the Office for Human Research Protections (OHRP) and the Food and Drug Administration (FDA) final rules on IRB registration requirements. Respondents for this information collection are institutions or organizations operating IRBs designated by an institution under an assurance of compliance approved for federalwide use by OHRP under 45 CFR 46.103(a) and that review human subjects research conducted or supported by HHS, or, in the case of FDA's regulation, each IRB in the United States that reviews clinical investigations regulated by FDA under sections 505(i) or 520(g) of the Federal Food, Drug and Cosmetic Act; and each IRB in the United States that reviews clinical investigations that are intended to support applications for research or marketing permits for FDA-regulated products.

ESTIMATED ANNUALIZED BURDEN TABLE

Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden hours
IRB Registration 0990–0279	6,000	2	1	12,000
FDA–IRB	1,000	2	1	2,000
Total				14,000

Terry Nicolosi,

Office of the Secretary, Paperwork Reduction Act Reports Clearance Officer.

[FR Doc. E9–6429 Filed 3–23–09; 8:45 am]

BILLING CODE 4150–36–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institute for Occupational Safety and Health; Decision To Evaluate a Petition To Designate a Class of Employees for the Baker Perkins Atomic Weapons Employer Facility in Saginaw, MI, To Be Included in the Special Exposure Cohort

AGENCY: National Institute for Occupational Safety and Health (NIOSH), Department of Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: HHS gives notice as required by 42 CFR 83.12(e) of a decision to evaluate a petition to designate a class of employees for the Baker Perkins Atomic Weapons Employer facility in Saginaw, Michigan, to be included in the Special Exposure Cohort under the Energy Employees Occupational Illness Compensation Program Act of 2000. The initial proposed definition for the class being evaluated, subject to revision as warranted by the evaluation, is as follows:

Facility: Baker Perkins Atomic Weapons Employer.

Location: Saginaw, Michigan.

Job Titles and/or Job Duties: All employees.

Period of Employment: May 14, 1956 through July 12, 1968.

FOR FURTHER INFORMATION CONTACT: Larry Elliott, Director, Office of Compensation Analysis and Support, National Institute for Occupational Safety and Health (NIOSH), 4676 Columbia Parkway, MS C–46, Cincinnati, OH 45226, Telephone 513–533–6800 (this is not a toll-free number). Information requests can also be submitted by e-mail to *OCAS@CDC.GOV*.

Christine M. Branche,

Acting Director, National Institute for Occupational Safety and Health.

[FR Doc. E9–6368 Filed 3–23–09; 8:45 am]

BILLING CODE 4163–19–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institute for Occupational Safety and Health; Decision To Evaluate a Petition To Designate a Class of Employees for the Electro-Metallurgical Corporation Facility, Niagara Falls, NY, To Be Included in the Special Exposure Cohort

AGENCY: National Institute for Occupational Safety and Health (NIOSH), Department of Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: HHS gives notice as required by 42 CFR 83.12(e) of a decision to evaluate a petition to designate a class of employees for the Electro-Metallurgical Corporation facility, Niagara Falls, New York, to be included in the Special Exposure Cohort under the Energy Employees Occupational Illness Compensation Program Act of 2000. The initial proposed definition for the class being evaluated, subject to revision as warranted by the evaluation, is as follows:

Facility: Electro-Metallurgical Corporation.

Location: Niagara Falls, New York.

Job Titles and/or Job Duties: All employees.

Period of Employment: August 13, 1942 through December 31, 1953.

FOR FURTHER INFORMATION CONTACT: Larry Elliott, Director, Office of Compensation Analysis and Support, National Institute for Occupational Safety and Health (NIOSH), 4676 Columbia Parkway, MS C–46, Cincinnati, OH 45226, Telephone 513–533–6800 (this is not a toll-free number). Information requests can also be submitted by e-mail to *OCAS@CDC.GOV*.

Christine M. Branche,

Acting Director, National Institute for Occupational Safety and Health.

[FR Doc. E9–6380 Filed 3–23–09; 8:45 am]

BILLING CODE 4163–19–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Annual Report/ACF 204 (State MOE)—1 collection.

OMB No.: 0970–0248.

Description: The Administration for Children and Families (ACF) is requesting a three-year extension of the ACF–204 (Annual MOE Report). The report is used to collect descriptive program characteristics information on the programs operated by States and Territories in association with their Temporary Assistance for Needy Families (TANF) programs. All State and Territory expenditures claimed toward States and Territories MOE requirements must be appropriate, i.e., meet all applicable MOE requirements. The Annual MOE Report provides the ability to learn about and to monitor the nature of State and Territory expenditures used to meet States and Territories MOE requirements, and it is an important source of information about the different ways that States and Territories are using their resources to help families attain and maintain self-sufficiency. In addition, the report is used to obtain State and Territory program characteristics for ACFs annual report to Congress, and the report serves as a useful resource to use in Congressional hearings about how TANF programs are evolving, in assessing State the Territory MOE expenditures, and in assessing the need for legislative changes.

Respondents: The 50 States of the United States, the District of Columbia, Guam, Puerto Rico, and the Virgin Islands.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
ACF–204	54	1	118	6,372
OLDC system updates	54	2	0.13	13.50

Estimated Total Annual Burden Hours: 6,385.50.

Additional Information

Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Administration, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. All requests should be identified by the title of the information collection. E-mail address: *infocollection@acf.hhs.gov.*

OMB Comment

OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, Fax: 202–395–6974, Attn: Desk Officer for the Administration for Children and Families.

Dated: March 19, 2009.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. E9–6452 Filed 3–23–09; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2008–N–0354]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Mental Models Study of Farmers' Understanding and Implementation of Good Agricultural Practices

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Fax written comments on the collection of information by April 23, 2009.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be faxed to the Office of Information and Regulatory Affairs, OMB, Attn: FDA Desk Officer, FAX: 202–395–6974, or e-mailed to *oira_submission@omb.eop.gov*. All comments should be identified with the OMB control number 0910–NEW and title, ''Mental Models Study of Farmers' Understanding and Implementation of Good Agricultural Practices.'' Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Jonna Capezzuto, Office of Information Management (HFA–710), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–796–3794.

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Mental Models Study of Farmers' Understanding and Implementation of Good Agricultural Practices

The proposed information collection will help FDA protect the public from foodborne illness by increasing the agency's understanding of how farmers and growers use Good Agricultural Practices (GAPs) to address common risk factors in their operations and thereby minimize food safety hazards potentially associated with fresh produce. Fresh fruits and vegetables are those that are likely to be sold to consumers in an unprocessed or minimally processed (i.e., raw) form and that are reasonably likely to be consumed raw. Under section 903(b)(2) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 393 (b)(2)), FDA is authorized to conduct research relating to foods and to conduct educational and public information programs relating to the safety of the Nation's food supply. Under Title 42 of the Public Health Service Act (1944), FDA has authority to act to protect the public health.

In 1998, FDA issued a guidance document entitled ''Guide to Minimize